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FAN.CNT 1

PATENT NO.

KIND DATE

(FILE 'HOME' ENTERED AT 19:20:45 ON 02 DEC 2003) FILE 'MEDLINE, CAPLUS, BIOSIS, SCISEARCH' ENTERED AT 19:20:58 ON 02 DEC L1846429 S PLASMID OR VECTOR L2 115648 S (CITRIC OR TARTARIC) (W) ACID L3 20 S L1(10A)L2 L417 DUP REM L3 (3 DUPLICATES REMOVED) => D BIB AB 1-17 L4 T.4 ANSWER 1 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN ΑN 2003:717784 CAPLUS DN139:231487 TΙ High flow compositions of compatibilized poly(arylene ether)-polyamide blends containing dendritic polyesters IN Adedeji, Adeyinka PΑ USA SO U.S. Pat. Appl. Publ., 8 pp. CODEN: USXXCO DTPatent LΑ English FAN.CNT 1 PATENT NO. KIND DATE APPLICATION NO. DATE _____ ----______ A1 20030911 A1 20030925 20030911 PΙ US 2003171503 US 2002-683955 20020306 WO 2003078526 WO 2002-US24000 20020708 AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG TJ, TM PRAI US 2002-683955 20020306 Α A thermoplastic compn. comprises a compatibilized poly(arylene ether)/polyamide resin blend and a dendritic polyester resin. A compatibilizer is a polycarboxylic acid, such as citric acid, and the compn. can further comprise an impact modifier, such as styrene-butadiene-styrene (SBS) block copolymer. Thus, a polyoxyphenylene (47.0), a polyamide (Capron 1250) (41.3), SBS block copolymer (Vector 8508D) (10.0), citric acid (0.8%), and a dendritic polyester (Boltorn H 20) were extruded and pelletized. compn. contg. 4.0% of the dendritic polyester had a melt flow rate of 16.33 (ASTM D1238) compared to 0.78 for a dendritic polyester-free blend, the rates being measured at the same conditions. ANSWER 2 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN L4ΑN 2002:84321 CAPLUS DN 136:147477 TI Method and kit for measuring tartaric acid-resistant acid phosphatase IN Miyazaki, Shuichi; Igarashi, Makoto PA Yamasa Shoyu Co., Ltd., Japan Jpn. Kokai Tokkyo Koho, 7 pp. SO CODEN: JKXXAF DTPatent Japanese LA

APPLICATION NO. DATE

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PI JP 2002031640 A2 20020131
PRAI JP 2000-216167 20000717
                                                   JP 2000-216167 20000717
      A method and a kit are provided for measuring tartaric acid-resistant acid
      phosphatase (TRAP) in a sample by an immunoassay. In this immunoassay,
      the antibody obtained by immunization using osteosarcoma cell-derived
      recombinant TRAP is used as an antibody, and the osteosarcoma cell-derived
      recombinant TRAP is used as a std. substance.
T<sub>1</sub>4
      ANSWER 3 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN
ΝA
      2002:609561 CAPLUS
DN
      137:151095
TI
      Preparation of supercoiled plasmid DNA by culture of bacteria in a defined
      medium
IN
      Voss, Carsten
PΑ
      Plasmidfactory Gmbh & Co. Kg, Germany
SO
      Ger. Offen., 22 pp.
      CODEN: GWXXBX
DT
      Patent
LA
      German
FAN.CNT 1
                                 DATE APPLICATION NO. DATE
      PATENT NO. KIND DATE
                          ____
      ______
      DE 10106493 A1 20020814 DE 2001-10106493 20010213 WO 2002064752 A1 20020822 WO 2002-EP290 20020114
PΙ
          W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GO, GW, MI, MR, NE, SN, TD, TG
               BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI DE 2001-10106493 A 20010213
      The present invention concerns a procedure for the prodn. of nucleic
      acids, esp. supercoiled DNA. The method involves cultivating a bacterial
      host carrying the plasmid to high cell densities in a batch process in a
      defined synthetic aq. medium that is free of complex components such as
      animal exts. The medium contains an org. carbon source, an inorg.
      nitrogen source, mineral salts, and an org. nitrogen compd. that supports
      bacterial metab., e.g. vitamins or amino acids. The purified nucleic
      acid, isolated from bacteria cells is suitable for use in gene therapy,
      cell therapy or genetic inoculation. Optimization expts. in which the
      effect of medium compn. and fermn. conditions on increasing the yield of
      the plasmid are described.
     ANSWER 4 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN DUPLICATE 1
L_4
AN
      2002:689576 CAPLUS
DN
      138:88727
      Enhancement of citric acid production by immobilized and freely suspended
      Aspergillus niger using silicone oil
ΑU
      Ates, Selma; Dingil, Nesrin; Bayraktar, Emine; Mehmetoglu, Ulku
CS
      Faculty of Art and Science, Department of Chemistry, Gazi University,
      Teknikokullar, Ankara, 06500, Turk.
      Process Biochemistry (Oxford, United Kingdom) (2002), 38(3), 433-436
SO
      CODEN: PBCHE5; ISSN: 1359-5113
PΒ
     Elsevier Science Ltd.
DT
     Journal
LΑ
     English
     The use of silicone oil as an oxygen vector for increasing
      citric acid prodn. by free and immobilized Aspergillus
      niger conidiospores was studied. When silicone oil was used, citric acid
      concn. increased with respect to the control run in free and immobilized
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systems 2.0 and 1.6 times, resp. The effect of potassium ferrocyanide [K4Fe(CN)6] on citric acid prodn. in a medium contg. air, oxygen and silicone oil was studied. When K4Fe(CN)6 was used with oxygen and silicone oil, citric acid concn. decreased significantly because of the conversion of ferrocyanide to ferricyanide. The reuse of immobilized A. niger conidiospores was investigated in citric acid prodn. medium contg. 2% (vol./vol.) silicone oil. Citric acid prodn. decreased on increasing the reuse no.

RE.CNT 30 THERE ARE 30 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L4 ANSWER 5 OF 17 SCISEARCH COPYRIGHT 2003 THOMSON ISI on STN
- AN 2003:22710 SCISEARCH
- GA The Genuine Article (R) Number: 622UL
- TI Enhancement of citric acid production by immobilized and freely suspended Aspergillus niger using silicone oil
- AU Ates S (Reprint); Dingil N; Bayraktar E; Mehmetoglu U
- CS Gazi Univ, Fac Art & Sci, Dept Chem, TR-06500 Ankara, Turkey (Reprint); Ankara Univ, Fac Engn, Dept Chem Engn, TR-06100 Ankara, Turkey
- CYA Turkey
- SO PROCESS BIOCHEMISTRY, (NOV 2002) Vol. 38, No. 3, pp. 433-436.
 Publisher: ELSEVIER SCI LTD, THE BOULEVARD, LANGFORD LANE, KIDLINGTON,
 OXFORD OX5 1GB, OXON, ENGLAND.
 ISSN: 0032-9592.
- DT Article; Journal
- LA English
- REC Reference Count: 30
 - *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS*
- The use of silicone oil as an oxygen vector for increasing citric acid production by free and immobilized Aspergillus niger conidiospores was studied. When silicone oil was used, citric acid concentration increased with respect to the control run in free and immobilized systems 2.0 and 1.6 times, respectively. The effect of potassium ferrocyanide [K4Fe(CN)(6)] on citric acid production in a medium containing air, oxygen and silicone oil was studied. When K4Fe(CN)(6) was used with oxygen and silicone oil, citric acid concentration decreased significantly because of the conversion of ferrocyanide to ferricyanide. The reuse of immobilized A. niger conidiospores was investigated in citric acid production medium containing 2% (v/v) silicone oil. Citric acid production decreased on increasing the reuse number. (C) 2002 Elsevier Science Ltd. All rights reserved.
- L4 ANSWER 6 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN
- AN 2001:578309 CAPLUS
- DN 136:101411
- TI Plasmid profile and characterization on negative mutants for lactose and citrate metabolism derived from Leuconostoc mesenteroides
- AU Sewaki, Tomomitsu; Tagawa, Yuji; Miyamoto, Taku
- CS Fac. Agric., Okayama Univ., Okayama-shi, 700-8530, Japan
- SO Miruku Saiensu (2001), 50(2), 49-54 CODEN: MISAFD; ISSN: 1343-0289
- PB Nippon Rakuno Kagakkai
- DT Journal
- LA Japanese
- AB Lactose- and citrate-neg. (Lac- and Cit-) mutants were isolated after the treatment of Leuconostoc mesenteroides strains 6-1-9 and OR-2 with acridine orange and examd. for their plasmid profiles and enzymic characteristics. Lac- mutants, designated 6-1-9-1 and 6-1-9-2 were deficient a 38 Mdal plasmid and they lost activities of the lactose-splitting enzyme (.beta.-galactosidase). On the other hand Citmutant (OR-2-1) missing a 15 Mdal plasmid lost the citrate permease activity, although it possessed less citrase activity.
- L4 ANSWER 7 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN

- AN 2000:750280 CAPLUS
- DN 133:307122
- TI Alcaligenes cis-epoxysuccinic acid hydrolase genes, and use in D-(-)-Tartaric acid production
- IN Asai, Yoko; Kobayashi, Tsuyoshi; Uchida, Koichi; Terasawa, Masato
- PA Mitsubishi Chemical Corp., Japan
- SO Jpn. Kokai Tokkyo Koho, 18 pp. CODEN: JKXXAF
- DT Patent
- LA Japanese

FAN.CNT 1

PATENT NO. KIND DATE APPLICATION NO. DATE
PI JP 2000295992 A2 20001024 JP 1999-105827 19990413
PRAI JP 1999-105827 19990413

AB Cis-epoxysuccinic acid hydrolase isolated from Alcaligenes, its genes, recombinant expression, and use in prodn. of D-(-)-Tartaric acid, are disclosed. Genes coding for .alpha. and .beta. subunits of cis-epoxysuccinic acid hydrolase were isolated from Alcaligenes MCI3611 strain. Hydrolysis of cis-epoxysuccinic acid to D-(-)-Tartaric acid was obsd. in E. coli transformed with the cloned genes.

- L4 ANSWER 8 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN DUPLICATE 2
- AN 2000:521129 CAPLUS
- DN 133:206842
- TI Enhancement of citric acid production by Aspergillus niger using n-dodecane as an oxygen-vector
- AU Wang, Jianlong
- CS State Key Joint Laboratory of Environment Simulation and Pollution Control, Department of Environmental Science and Engineering, Tsinghua University, Beijing, 100084, Peop. Rep. China
- SO Process Biochemistry (Oxford) (2000), 35(10), 1079-1083 CODEN: PBCHE5; ISSN: 1359-5113
- PB Elsevier Science Ireland Ltd.
- DT Journal
- LA English
- AB Due to the significant oxygen requirement during citric acid prodn. and the relatively low soly. of oxygen in water, aeration is crit. The potential use of n-dodecane as an oxygen-vector for improvement of citric acid prodn. by Aspergillus niger was studied. The volumetric fraction of oxygen-vector has a great influence on the volumetric oxygen transfer coeff. kLa. With the addn. of an oxygen-vector to the fermn. medium with a final concn. of 5%, the kLa value reached a max. value (130 h-1), which is twice that of the control expt. The addn. of 5% (vol./vol.) n-dodecane enhanced citric acid accumulation, reduced residual sugar concn. and stimulated mycelial growth. Adding n-dodecane had no adverse effects on the cells of A. niger. The results of enzyme assays indicated that no significant differences were obsd. between the activity of citrate synthase of two kinds of mycelial cell-free exts.
- RE.CNT 18 THERE ARE 18 CITED REFERENCES AVAILABLE FOR THIS RECORD ALL CITATIONS AVAILABLE IN THE RE FORMAT
- L4 ANSWER 9 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN
- AN 2000:156429 CAPLUS
- DN 133:130432
- TI A biofunctional assay to study pRL-CMV plasmid DNA formulation stability
- AU Poxon, Scott W.; Hughes, Jeffrey A.
- CS Department of Pharmaceutics, College of Pharmacy, University of Florida, Gainesville, FL, 32610, USA
- SO PDA Journal of Pharmaceutical Science and Technology (1999), 53(6), 314-317
 - CODEN: JPHTEU; ISSN: 1076-397X
- PB PDA, Inc.

- DT Journal
- LA English
- The ability of a plasmid DNA formulation to code for a functional protein was assayed as a marker for plasmid DNA stability using a cotransfection method to measure transcription efficiency. This method shows increased sensitivity and reproducibility over single plasmid transfection methods. Method validation, by measuring DNA degrdn. rates, demonstrates that buffer choice may be of some importance in the pharmaceutical formulation of plasmid DNA. Degrdn. rates dependent on citrate buffer concn. were obsd. This cotransfection method has proven superior to std. agarose gel electrophoresis in quantifying subtle pRL-CMV plasmid DNA damage and could be used to help predict stability of a final plasmid DNA dosage form.

RE.CNT 12 THERE ARE 12 CITED REFERENCES AVAILABLE FOR THIS RECORD
ALL CITATIONS AVAILABLE IN THE RE FORMAT

- L4 ANSWER 10 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN
- AN 1994:291318 CAPLUS
- DN 120:291318
- TI Carbon source-dependent inhibition of xyl operon expression of the Pseudomonas putida TOL plasmid
- AU Holtel, Andreas; Marques, Silvia; Moehler, Isabel; Jakubzik, Ute; Timmis, Kenneth N.
- CS Dep. Microbiol., GBF-Natl. Res. Cent. Biotechnol., Braunschweig, Germany
- SO Journal of Bacteriology (1994), 176(6), 1773-6 CODEN: JOBAAY; ISSN: 0021-9193
- DT Journal
- LA English
- AB TOL plasmid-encoded degrdn. of benzyl alc. by Pseudomonas putida is inhibited by glucose and other compds. related to the main carbohydrate metab. in Pseudomonas species. The authors report here that this effect is exerted at the level of expression of the xyl catabolic operons, and two xyl promoters, Pu and Ps, were identified as the primary targets of this inhibition. Xyl promoter activation was also inhibited by glucose in the heterologous Escherichia coli system, apparently not however by the classical mechanism of enteric catabolite repression.
- L4 ANSWER 11 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN
- AN 1991:181893 CAPLUS
- DN 114:181893
- TI Characterization of a citrate-negative mutant of Leuconostoc mesenteroides subsp. mesenteroides: metabolic and plasmidic properties
- AU Lin, J.; Schmitt, P.; Divies, C.
- CS Dep. Microbiol. Biotechnol., Ec. Natl. Super. Biol. Appl. Nutr. Aliment., Dijon, F-21000, Fr.
- SO Applied Microbiology and Biotechnology (1991), 34(5), 628-31 CODEN: AMBIDG; ISSN: 0175-7598
- DT Journal
- LA English
- AB Comparison of the parental strain of the L. mesenteroides subsp.

 mesenteroides (19D) and its citrate-neg. mutant, which has lost a 22-kb plasmid, has confirmed the energetic role of citrate. Fermn. balance anal. showed that citrate led to a change in heterolactic fermn. from glucose. High levels of enzyme activity in both mutant and parental strains were found for NADH oxidase, lactate dehydrogenase, acetate kinase, alc. dehydrogenase, diacetyl reductase and acetoin reductase, although NADH oxidase, alc. dehydrogenase, diacetyl reductase, and acetoin reductase were partly repressed by citrate. All these enzymes studied were not plasmid-linked. In the parental strain, citrate lyase was induced by citrate. No citrate lyase activity was found in the citrate-neg. mutant grown in presence of citrate, but this does not provide evidence that citrate lyase is linked to the 22-kb plasmid.
- L4 ANSWER 12 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN
- AN 1991:510283 CAPLUS

- 115:110283 DN
- Instability of lactose and citrate metabolism of Leuconostoc strains TI
- Fantuzzi, L.; Vescovo, M.; Bottazzi, V. ΑU
- Ist. Microbiol., Univ. Cattol., Piacenza, 29100, Italy
 Biotechnology Letters (1991), 13(6), 433-6 CS
- CODEN: BILED3; ISSN: 0141-5492
- DT Journal
- LΑ English
- The instability of Lac+ and Cit+ phenotypes was investigated in Leuconostoc mesenteroides cremoris ATCC 19245 and in 4 strains of L. mesenteroides dextranicum. The 2 phenotypes were linked to a 14-Mdal and a 34-Mdal plasmid, resp., in L. mesenteroides cremoris ATCC 19245. In L. mesenteroides dextranicum, the character Lac+ was linked to a 28-Mdal plasmid, while the Cit+ phenotype was stable.
- ANSWER 13 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN DUPLICATE 3 L4
- 1989:495542 CAPLUS ΑN
- DN111:95542
- TIGlucose as a substrate in recombinant strain fermentation technology. By-product formation, degradation and intracellular accumulation of recombinant protein
- ΑÜ Rinas, Ursula; Kracke-Helm, Heinrich Andreas; Schuegerl, Karl
- Inst. Biophys. Phys. Biochem., Univ. Regensburg, Regensburg, D-8400, Fed. CS
- SO Applied Microbiology and Biotechnology (1989), 31(2), 163-7 CODEN: AMBIDG; ISSN: 0175-7598
- DTJournal
- English LA
- AΒ Glucose supplements to complex growth medium of Escherichia coli affect the prodn. of a recombinant model protein under the control of a temp.-sensitive expression system. The bacterial Crabtree effect, which occurs in the presence of glucose under aerobic conditions, not only represses the formation of citric acid cycle enzymes, but also represses the formation of the plasmid-encoded product, even though the synthesis of this protein is under the control of the temp.-inducible lambda PR promoter/cI857 repressor expression system. When the recombinant E. coli is grown at a moderate temp. (35.degree.) with protein hydrolyzate and glucose as substrates, a biphasic growth and prodn. pattern is obsd. In the 1st phase, the cells grow with a high specific growth rate, utilizing glucose and forming glutamate as a byproduct. The intracellular level of recombinant protein is very low in this phase. Later, glutamate is consumed, indicating an active citric acid cycle. The degrdn. of glutamate is accompanied by the intracellular accumulation of high amts. of recombinant protein.
- ANSWER 14 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN L4
- 1989:113201 CAPLUS AN
- DN110:113201
- L-Glutamic acid and L-proline, their recombinant manufacture with Corynebacterium and Brevibacterium
- IN Katsumata, Ryoichi; Yokoi, Haruhiko; Kino, Kuniki
- PAKyowa Hakko Kogyo Co., Ltd., Japan
- SO Jpn. Kokai Tokkyo Koho, 16 pp. CODEN: JKXXAF
- DT Patent
- Japanese
- FAN.CNT 1

	PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
ΡI	JP 63119688	A2	19880524	JP 1986-265297	19861107
	JP 07121228	B4	19951225		
PRAI	JP 1986-265297		19861107		

AΒ Glutamic acid and proline are manufd. by cultivating recombinant Corynebacterium or Brevibacterium contg. the gene encoding citric acid synthase. Plasmid pEgltA-1 contg. the synthase gene cloned from the chromosomal DNA of Escherichia coli was linked to plasmid pCG11, a vector for both Corynecbacterium and Brevibacterium, to form recombinant plasmid pEgltA-2. Corynebacterium glutamicum transformed with pEgltA-2 produced glutamic acid 31.2 mg/mL culture fluid.

- L4 ANSWER 15 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN
- AN 1983:483242 CAPLUS
- DN 99:83242
- TI Bacterial-plant gene cloning shuttle vectors for genetic modification of plants
- AU Kado, C. I.; Tait, R. C.
- CS Dep. Plant Pathol., Univ. California, Davis, CA, 95616, USA
- SO NATO ASI Series, Series A: Life Sciences (1983), 61(Genet. Eng. Eukaryotes), 103-10
 CODEN: NALSDJ; ISSN: 0258-1213
- DT Journal
- LA English
- AB A discussion of the construction of plasmid cloning vectors from Agrobacterium tumefaciens, which can potentially shuttle genes between Escherichia coli and plants, is presented. These include plasmid derivs. of pTAR and pSa. The pTAR plasmids contain the genetic determinants for the stereospecific catabolism of L-tartaric acid in Agrobacterium. Plasmid pCK2G, a pTAR deriv., contains the origin of replication of pTAR and the E. coli plasmid pBR322 and can be introduced into either E. coli or A. tumefaciens by transformation. Plasmid pTi and pCK2G are both compatible. DNA sequences present on both plasmids can be exchanged by homologous recombination. Thus, a pCK2G recombinant contg. a fragment of T-DNA is a useful cloning vector for plant genes. Plasmid pSa151-T utilizes the origin of replication of the S strain of the cauliflower mosaic virus genome in the PvuII site of pSa151 and can replicate in plant hosts.
- L4 ANSWER 16 OF 17 CAPLUS COPYRIGHT 2003 ACS on STN
- AN 1982:156615 CAPLUS
- DN 96:156615
- TI Genetic and molecular studies of the regulation of atypical citrate utilization and variable Vi antigen expression in enteric bacteria
- AU Baron, L. S.; Kopecko, D. J.; McCowen, S. M.; Snellings, N. J.; Johnson, E. M.; Reid, W. C.; Life, C. A.
- CS Dep. Bacterial Immunol., Walter Reed Army Inst. Res., Washington, DC, 20012, USA
- SO Basic Life Sciences (1982), 19, 175-94 CODEN: BLFSBY; ISSN: 0090-5542
- DT Journal
- LA English
- Atypical citric acid [77-92-9] utilization by AΒ Escherichia coli strains V414 and V517 was plasmid-encoded, by a 130-megadalton conjugative Cit+ plasmid, (pWR60) in V414 and by a 36-megadalton plasmid (pWR517-7) in V517. The Cit+ genes of pWR60, present on a 9-kilobase PstI fragment, were cloned in pBR325. Atypical citrate utilization apparently involved partial metab. of citrate at the cell surface, before or during uptake. Expression of atypical citrate utilization encoded by pWR60 or a recombinant (pWR61) appeared to be reversible. The genes viaA and viaB were necessary for Vi antigen formation by Salmonella and Citrobacter. Reversible expression of the Vi antigen by some C. freundii strains was controlled by the viaB locus, which also encoded the Vi antigen. S. typhi And E. coli K12 hybrid strains carrying the C. freundii viaB locus exhibited reversible Vi antigen expression, even in the absence of general recombination. The viaB locus of C. freundii was transferred to an F'lac plasmid in E. coli K12 strains WR2376 via phage Mu-mediated transposition.

- AN 1980:87128 BIOSIS
- DN PREV198019024626; BR19:24626
- TI GENETIC CHARACTERISTICS OF CITRIC-ACID UTILIZING PLASMID COEXISTING WITH H-1 GROUP R PLASMID IN SALMONELLA-TYPHIMURIUM OF COW ORIGIN.
- AU SATO G [Reprint author]; ISHIKURO N; OKA C; ASAKI M; HANZAWA Y
- CS DEP VET MED HYG, OBIHIRO VET COLL, OBIHIRO, HOKKAIDO, JPN
- Japanese Journal of Bacteriology, (1979) Vol. 34, No. 1, pp. 149.
 Meeting Info.: 52ND MEETING OF THE JAPANESE SOCIETY FOR BACTERIOLOGY,
 HIRATSUKA CITY, JAPAN, APR. 4-6, 1979. JPN J BACTERIOL.
 CODEN: NSKZAM. ISSN: 0021-4930.
- DT Conference; (Meeting)
 Conference; Abstract; (Meeting Abstract)
- FS BR

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LA JAPANESE